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Gary Paulazzo

**MECHANICAL CLOT TREATMENT DEVICE  
WITH DISTAL FILTER**

CROSS REFERENCE TO RELATED PATENT APPLICATION

5 This application is a continuation of U.S. Patent Application Serial No. 08/941,514, filed September 30, 1997, the entirety of which is hereby incorporated by reference.

FIELD OF THE INVENTION

10 This is a medical device used to treat embolic strokes. In particular, it is a surgical device usually delivered through an intravascular catheter. It may be used in several ways. It may, for instance, be used to open a clear passageway adjacent a thrombus to allow both blood and medication to bypass the clot. It may be used to pierce and to remove a thrombus. These thrombi are often found in tortuous vasculature. The device may include several sections: an elongated core element, a proximal cage assembly, a distal filter, and an actuator. The core element is at least a

15 core wire. Placed around the distal end of the core element is a collapsible, but preferably self-expanding or self-collapsing, proximal cage assembly and a more distally located, preferably self-expanding or self-collapsing, distal filter, cage assembly, or other filter component. These components are preferably radio-opaque. The proximal end of the proximal cage is typically is affixed to an actuator in such a

20 way as to allow expansion of the cage after deployment. The proximal cage assembly may be used for collecting emboli or for displacing them to allow blood flow to resume, either with or without concurrent clot-dissolving drug treatment. The distal

sector, whether a cage or a fan, is placed distally of the thrombus to collect portions of the thrombus which may loosen during the treatment or removal procedure.

### BACKGROUND OF THE INVENTION

5           This surgical device is designed to displace or to penetrate emboli found in the human vasculature. The device has several major components including a proximal (or displacing cage) and a distal filter. Depending upon the chosen procedure, the device is inserted so that the displacing or proximal cage is either towards the venous side of the clot or adjacent the clot. The proximal cage is expanded either to displace  
10           the clot away from the arterial wall and allow flow of fluid (e.g., blood and medications such as anti-thrombolytics or other lysing agents) past (or to) the formerly occluded site or to engage the clot for removal using the cage as a collector assembly, expand once past the target emboli, and catch or net the embolism (or a portion of the embolism) for removal from patient's blood vessels. In some  
15           situations, the device may be used to move the clot to another position in the vasculature; perhaps for recovery using another device or to canalize the clot for improved blood flow. The distal filter member is always placed distal to the thrombus to catch any portions which may break away.

          Other devices to treat vascular emboli are known. The use of inflatable  
20           balloons to remove emboli has been practiced for many years. The "Fogarty catheter" has been used, typically in the periphery, to remove clots from arteries found in legs and in arms. These well known devices have been described in some detail in U.S. Patent No. 3,435,826, to Fogarty and in U.S. Patent Nos. 4,403,612 and 3,367,101. These patents describe a balloon catheter in which a balloon material is  
25           longitudinally stretched when deflated.

          Emboli occasionally form around the valves of the heart and then are dislodged and follow the blood flow into the distal regions of the body. They are

particularly dangerous if the emboli passes to the brain and causes an embolic stroke. As will be discussed below, many such occlusions lodge in the middle cerebral artery (MCA), although such is not the only site where emboli come to rest. Obviously, when blood flow is inhibited or cut off completely from a portion of the brain, the brain's oxygen supply is limited causing severe health problems.

In procedures for removing emboli using the Fogarty catheter or other similar catheters, it is typical, first, to locate the clot using fluoroscopy. The embolectomy catheter is then inserted and directed to the clot. The distal tip of the balloon catheter is then carefully moved through the center of the clot. Once the balloon has passed through the distal side of the clot, the balloon is inflated. The balloon catheter is then gradually and gently withdrawn. The balloon, in this way, acts to pull the clot ahead of the balloon. The majority of procedures using a Fogarty catheter repeat these steps until the pertinent vessel is cleared of clot material.

Such vaso-occlusions occur in a wide variety of sites within the body. The lodging of thrombus in various sites is often complicated by the presence of atherosclerosis. This disease causes the vessels to become tortuous and narrowed. These anomalies are often considered to be the result of the growth of atherosclerotic plaque. Clots occurring in these diseased vessels are difficult to remove using balloon or Fogarty catheters.

Removal of emboli using balloon catheters is rife with potential problems. One such problem occurs during removal of a clot. The resistance to such removal often causes the balloon portion of the catheter to evert over the tip of the catheter. Should the user need to partially deflate the balloon during such a deflation, the distal tip of the balloon may become distended and angulate. Another difficulty with balloon catheters is the possibility of damage to the intima of arteries. Inflation pressures can create forces significant enough to score such a vessel lining or

dislodge plaque lodged on such a wall. In the worst case, the balloon may rupture leaving balloon portions in the bloodstream.

5 Movement of a balloon in the MCA using only a balloon can displace the clot through more proximal branches into other large vessels such as the internal carotid artery (ICA) and then into other vessels and may pressure the clot into branching vessels.

There are a variety of different devices intended for use in replacing balloon catheters and in using a device other than a balloon catheter in so removing the emboli.

10 One such device is shown in U.S. Patent No. 4,030,503 to Clark III. This patent describes a spiral helix affixed to the distal end of a catheter. In particular, the spiral helix is designed to be rotated and pushed forward through the clot. It is said that the helix screws into the clot, and when it is firmly embedded or is past the clot, the catheter is pulled out of the vessel without rotation. The catheter is said to operate  
15 like a corkscrew.

A similar catheter is described in U.S. Patent No. 4,706,671 to Weinrib. This catheter also has a coil section at its distal end. The coil section is said to be stretched initially into a generally linear insertion position for removing inwardly in a vessel. The coil member is then expanded into the form of a hollow conical scoop to then  
20 scoop clot material from the blood vessel. The coil member is stiffened by an internal wire which is then removed. The hollow passageway is then filled with a liquid to stiffen the coils. The coils are said to be of an elastomeric material.

U.S. No. Patent 4,762,130 to Fogarty et al., describes a helical balloon attached to the distal end of a catheter. The helical or bellowed balloon is maintained  
25 in a generally linear condition and passed into a clot. Once the catheter balloon within the clot is inflated, the balloon and adjoining clot are removed together.

Another similar device used more to grip and shear atherosclerotic deposits rather than to remove thrombi is described in U.S. Patent No. 4,890,611 to Monfort et al. This device incorporates a pair of helical wires placed on the distal end of a wire. The flexible wire is pulled against a flexible catheter and the two helically  
5 configured loops expand to form a shearing apparatus. The totality of the apparatus is then twisted by means of a handle so that the pair of helically wound loops cuts through and is said to retain sections of plaque for removal from the vessel.

Another thrombus extraction system is shown in U.S. Patent No. 5,011,488, to Ginsberg. In using this device, an inflatable balloon having a proximal conic shape is  
10 deflated and passed through a thrombus. It is then expanded and retracted so that the proximal passage pulls the thrombus into contact with an aspirator. The aspirator then removes the clot or thrombotic material from the vessel.

An alternative configuration of the expandable member is also described in the Ginsberg patent mentioned just above. In this variation, a wire coil is attached to  
15 an extension wire which may be moved between an extended position and a retracted position. The retracted or expanded configuration is illustrated to have a conical shape. The cone is shown to be one which has a smaller end proximally.

U.S. Patent No. 5,112,347, to Taheri, shows an inflatable balloon type embolectomy catheter. The balloon has a number of fingers arranged in a leaf spring  
20 arrangement inside the balloon. The balloon is hydraulically inflated and forms a cone after inflation. The deflated device is shown in Figures 11 through 14 to be passed distally past an embolism before inflation. After inflation, the large end of the balloon collects the embolism as it is pulled past the appropriate site in the vessel.

U.S. Patent No. 5,192,286, to Phan, shows a retrieval catheter for removing  
25 materials from various body lumen. The retrieval catheter is shown to have a slack net which may be collapsed for passage into a lumen past the material to be collected. The net is unfolded and materials such as uretral stones are removed.

U.S. Patent No. 5,411,509 to Hilal, shows an embolectomy catheter having an elastomeric foam tip attached distally. The foam tip has an actuator means suitable for forming the foam section both longitudinally and radially in response to activation of the actuation means. In practice, the catheter tip is pressed past an embolism, inflated, and retracted with the clot being pushed proximally as retraction occurs.

U.S. Patent No. 5,490,859, to Mische et al., shows an intravascular occlusion material removal device having an expandable material removal element made up of a number of wires passing between the two ends of such element, a catheter shaft, a drive shaft for spinning the material movement element within the blood vessel, and a collection portion placed on the material removal element for collecting any occlusion material removed by the expandable material removal element. The drive shaft may be operated by a motor connected to the proximal end of the drive shaft.

None of these devices show the device described below.

#### SUMMARY OF THE INVENTION

This is a surgical device usually delivered through an intravascular catheter. It is designed variously to displace or to pierce and remove emboli particularly when found in tortuous vasculature.

This embolectomy device includes several sections. First, the device has a core element. The core element preferably has at least several major functions: first, the distal portion may act as a guidewire for vessel navigation to the treatment site; second, the distal portion assists in puncturing the clot or displacing it away from the arterial wall; and third, it cooperates with the collection assembly during the deployment of that assembly. The core element may be a simple core wire (perhaps enhanced with an adherent braid) fixedly attached to the distal end of the cage assembly or a tubular member attached to the collection assembly having a removable core or guidewire in its interior passageway. Preferably, the core element is able to

rotate freely with respect to collection assembly to preserve its abilities as a guidewire while preserving the cage assembly shape and integrity.

Placed distally on the core element are a pair of cage or filter elements. The proximal element preferably is an expandable and collapsible self expanding cage assembly. The proximal element may be self-expanding or self-collapsing. The distal element may also be a cage and design similar to the proximal element or of other filter design. The distal element is intended to catch any errant thrombus or clot fragments. Both elements may either be fixedly or rotatably joined to the core wire. Most preferred of the designs for the cage assembly involves a super-elastic alloy ribbon or wire braid wrapped with a platinum wire or otherwise made radio-opaque.

The assembly further may have an actuator which permits or causes the collector assembly to expand or to contract after deployment. The actuator may be mechanical in nature or electrolytic, or may include a balloon. When the actuator is mechanical, it is typically attached to the proximal end of the proximal cage assembly to allow or cause controllable expansion of the distal and proximal cage assemblies.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows a generalized schematic of a portion of the arterial system of the head specifying in particular the position of an embolism in the middle cerebral artery.

Figures 2A and 2B show a side, partial cutaway and a side view of the distal end of one variation of the device respectively as expanded and as unexpanded.

Figures 3A and 3B show, respectively, a side, partial cutaway and an end view of another variation of the device.

Figures 4A, 4B, and 4C show, respectively, side views of various distal tips of the device.

Figure 5A shows a close-up of braid section comprised of a wrapped wire.

Figure 5B shows a close-up of braid section comprised of a wrapped ribbon.

Figures 6A, 6B, 6C, and 6D show a generalized sequence of steps for use of the inventive device for displacing a thrombus for restoration of flow past the occluded site.

### DESCRIPTION OF THE INVENTION

This device is a surgical implement. It is designed variously to move a thrombus aside and provide a fluid passageway along that thrombus or to penetrate and at least partially to retrieve emboli situated in human vasculature. It is intended to be used rapidly as a flow restoration device. Even in those instances where the embolism is not or cannot be completely removed, this inventive device is believed to be useful in displacing, extracting, or removing a portion of the clot and thereby permitting restoration of partial blood flow. It can be used to move the clot to a more advantageous site in the body. It is suitably flexible to be placed in the distal tortuous vasculature of the brain and hence is useful in treating blocking emboli found there. This device may be considered a partial treatment for embolic stroke, for instance, when used with the addition of thrombolytics.

As a matter of practical experience, a large proportion of emboli sloughed into the brain are seldom longer than about 1 to 2 centimeters and, for a variety of reasons both physiological and hemodynamic, settle in the middle cerebral artery (MCA). As is shown in Figure 1, the path from the site of origin of a traveling embolus -- often the heart -- to the MCA is via the common carotid artery past the branch point forming the external and internal carotid arteries into the internal carotid artery (ICA). The MCA is generally considered to be the continuation and termination of the ICA after the siphon and after the branching sites of a variety of other arteries, e.g., the



ophthalmic artery, the anterior communicating artery, the posterior communicating artery, and others. The etiology of such an occlusion is varied, varying, and complicated. The occlusion (100) is shown in the MCA in Figure 1 at the noted site.

5        Treatments for such embolic occlusions include catheterization of the patient and introduction of tissue plasminogen activator (TPA) or urokinase or other clot lysing agents to the site of the occlusion. Additionally the embolic occlusion may be penetrated -- often with a microcatheter -- and the TPA or urokinase introduced distally of the occlusion. Removal of the catheter provides a modest passageway for resumed or increased blood flow past the then-partial occlusion.

10        This inventive device is for the rapid restoration of fluid or blood flow past the occlusion. We have found that it may be used in a variety of ways. For instance, when approaching the clot from the arterial end, the device can be used either to bypass the clot along the vessel wall and, upon expansion of the proximal cage assembly, to allow flow of blood across the occlusion using the interior of the  
15        proximal cage assembly as a fluid passageway.

      Alternatively, the inventive device may be used to canalize or remove an occlusion, via the steps of penetrating the occlusion, expanding a cage assembly distally of the embolism, and preferably removing at least a part of the embolism along with the cage and its attendant catheter.

20        The distal cage or filter element is to catch and retain any thrombus fragments and prevent them from moving distally.

      Figure 2A is a diagrammatic portrayal of the major portions of a desired variation (200) of the device and provides a general convention for several of the terms used herein.

25        The assembly typically includes a core element, often a core wire (202), and optional coil tip (204) to provide a measure of radio-opacity to the distal tip and protect against damage to the intima, a distal cage assembly, filter, or element (206)

which may be used as an embolism collector, and a proximal cage assembly or element (208). Because of the "fineness" of the device, the coil tip (204) may also serve the same function as a guidewire tip. The length of the coil tip (204) may be up to 4 or 5 centimeters in such instances. Further, and as is discussed below, the core element may be an assembly comprising a tubular member further containing a removable core or guidewire. In one highly desirable variation of the invention, a separation member (210) is used to separate distal cage or element (206) from the proximal cage (208). As will be explained in greater detail below, a preferred variation of the invention is one in which the distal member (206) and the proximal cage (208) are made of a single tubular member woven variously from wire or ribbon. The two cages are separated by a separation member (210) which, as seen in partial cross section in Figure 2A, is a simple tubular member made from, e.g., an appropriate metal or polymer.

The distal cage assembly (206) and proximal cage assembly (208) are collapsible for passage along-side of or for penetration and passage through the embolism and are generally self expandable once in position. Other components and devices may, obviously, be used to expand the distal cage assembly (206) and proximal cage assembly (208), are discussed at some length below.

The distal cage assembly (206) and proximal cage assembly may either self-expand when the actuator element (212) is permitted to move from its introduction position to its deployment position or it may self-collapse. Self-collapsing may provide a measure of safety in the event that the device becomes inoperable. Withdrawal of a collapsed device is simpler than is removal of an expanded device. The actuator element (212) is operable by the person.

Figure 2B shows the shape of the variation shown in Figure 2B in the form in which it is introduced. Actuator (212) (not shown) has been pulled proximally against the core wire (202) (not shown in Figure 2B). As will be shown below, this

slim or first configuration passes through an appropriately sized delivery catheter and may be pushed by or through a clot prior to expansion to the deployed or second configuration shown in Figure 2A.

5 The distal cage assembly (206) and proximal cage assembly (208) are preferably made of a super-elastic alloy wire or ribbon. Some stainless steels are suitable but the ready availability of nickel-titanium alloys in a wide variety of shapes and sizes makes this choice an easy one. In particular, we have found that ribbons as thin as 0.75 mils in thickness and as narrow as 2 mils in width are suitable for this device. Thicker and wider ribbons are also suitable in some instances as the situation  
10 requires. Preferred ribbons are between 0.75 and 1.5 mils in thickness and 3 and 7 mils in width. Most preferred are 0.8 mils and 4 mils respectively.

By "wires" we mean to include filamentary shapes having cross-sections where the thickness/width ratio is greater than 0.5 to and including 1.0. The cross-sectional form may be circular, square, oval, etc.

15 At least a minority of the ribbons may be a suitable stainless steel (e.g., 304SS, 306SS, etc.) or other such alloy. A desirable variant is the substitution of one or more ribbons of the braid (or the addition of one or more ribbons) with a radio-opaque material such as platinum. This obviously permits the user to visualize the position of the distal cage assembly (206) and proximal cage assembly (208) during a  
20 procedure.

A technical basis for the term "super-elastic" is found in the class of nickel-titanium alloys known as "nitinol" -- alloys discovered by the United States Naval Ordnance Laboratory. These materials are discussed at length in U.S. Patent  
25 Nos. 3,174,851 to Buehler et al., 3,351,463 to Rozner et al., and 3,753,700 to Harrison et al. Alloys especially suitable for this invention are those containing at least 1.5% (wt.) and up to about 8% (wt.) or more, of one or more alloying members selected from the group consisting of vanadium, chromium, manganese, iron, and

cobalt. By the term "ribbon", we intend to include elongated shapes, the cross section of which are not square or round and may typically be rectangular, oval, or semi-oval. They should have an aspect ratio of 0.5 (thickness/width) or less.

5 The distal cage (206) and proximal cage (208) may be made by winding a ribbon or a wire onto a mandrel having an exterior form to produce the relaxed shape found in Figure 2A. The mandrel and wound ribbon are preferably heated for a suitable period of time (e.g., 750° to 1200° F for one-half to two hours) to produce an assembly which has the portrayed shape. For use in the MCA, the outer diameter of the device after deployment need not be any larger than about 3 mm. It is desirable  
10 that the collapsed or non-deployed device have a diameter no larger than about 1 mm. The expansion ratio between any of the deployed and deployable devices discussed herein typically need not be any more than about 3:1 to be effective in the MCA. Expansion ratios between 2:1 and 10:1 are appropriate in this invention. These ratios are suitable for each variation of the invention. The invention is not so limited,  
15 however.

In some designs of the device, it is permissible to use an actuator as an active deployment element rather than as a mere restraint on the premature expansion of the cage assembly. Examples of such actuators are shown in U.S. Pat. Application  
20 Nos. 08/686,304, 08/701,155, and 08/869,346 assigned to Target Therapeutics and all of which are incorporated by reference.

Figures 3A and 3B show a cross-section of another variation of the inventive device related to that variation found in Figures 3A and 3B but employing a slightly more complicated distal cage or filter assembly.

25 Figures 3A and 3B show respectively a partial cross-section of a further variation (220) of the invention shown, as above, a device having a central core wire (222) and a proximal cage assembly (224). Proximal cage assembly (224) is shown to be a woven braid of ribbons, preferably of a superelastic alloy, but as with

variations in this Application, may be made of wires as that term is defined herein. It is to be seen from Figure 3A that the shape of the proximal cage assembly (224) is somewhat more columnar than that shown in Figure 2A. This shape is a preferred one because of its ease of manufacture. It may be produced in such a way that it is of significant length for use in long emboli. Actuator tubular member (226) is also shown in this variation, as is separator element (228). A major difference from the variation (200) shown in Figures 2A and 2B is the structure of the distal filter assembly (230). Although the structure of distal filter assembly (230) is not critical to the concepts of this invention in that it may be of any desirable shape capable of catching small bits of emboli sloughed from the embolus against which a proximal cage assembly (224) is pressing, the depicted distal filter assembly (230) is exemplative of the shape of assemblies suitable for placement on this section of the overall assembly. In this variation, a plurality of petal-shaped wires (232) which may or may not overlap when deployed, are seen. Optional spacer wires (234) are seen as supporting assembly tip member (236) to provide the distal end of core wire (238) with a measure of radial placement.

Figures 3A and 3B show the distal assembly in its deployed configuration. Because of the distal rake of filter members (232), those filter members (232) may be easily retracted into a catheter lumen when one desires to remove this device from the human body.

Figures 4A, 4B, and 4C show various distal treatments of the overall device. As has been noted above, the core wire may be fixed within the cage members, it may be provided with limited movement axially within the cage members, or it may be provided with full axial movement in which case the core wire acts and may be used as a guidewire for the device.

Figure 4A shows a very simple variation of the invention in which a core wire (242) is fixably attached to a distal coil (244). The distal coil (244) also serves as an

anchoring means for the distal cage assembly (246). This variation of the invention is simple and inexpensive and is suitably useful particularly when the device need not be extended too far into tortuous vasculature.

Figure 4B shows another variation of the invention also having a woven  
5 braided ribbon distal cage (248). However, unlike the variation shown in Figure 4A, this variation utilizes a core wire (250) which is freely moveable axially through the center of that distal cage (248). The distal cage (248) has a tubular liner (252) which moves independently of the actuator (shown in Figures 2A and 3A above). Core wire (250) does act as a guidewire and may be used to provide a pathway through tortuous  
10 vasculature for the device to follow. Liner (252) is preferably made of a material such as polytetrafluoroethylene (e.g., Teflon) and desirably extends all the way to the proximal end of the catheter and acts as an actuator for the cages. A desirable variation of this invention includes the securing of distal cage assembly (248) by use of a tubular member of, e.g., a thermoplastic, which fixes the distal-most end of distal  
15 cage assembly (248) to inner liner (252). A radio-opaque marker (256) (which is optional in this variation for reasons which will be explained below) is also depicted.

Figure 4C shows a further variation (258) of this invention in which core wire (260) is rotatable with respect to distal cage (262) but has limited axial movement therein. This obviously allows the user to position the bent coil tip (244) in a position  
20 to select or sub-select from turns or Y's in the tortuous vasculature without twisting the whole device (258). Limitation of the movement of the core wire (260) within the device (258) is accomplished via the use of bushing (264) placed distally in the distal cage assembly (262). The inner diameter of bushing (264) will not allow coil (244) to pass proximally nor will it allow coil (266) to pass distally. Coil (266) may be  
25 used as a marker coil if so desired. Bushing (264) also desirably forms a joint with outer tubular member (286) to firmly place the distal-most end of distal cage assembly (262).

An independent feature shown in Figure 4C is the presence of a braid member (268) which is adherent to core wire (260). Braid member (268) is for the specific purpose of providing kink resistance and improved torqueability. The material used as a braid on the core wire (260) may be any of a different number of materials.

5 Highly desirable, though, is the use of a braid of super-elastic alloy. Certain classes of these materials have been discussed in detail above. The braid material (268) may be stainless steel wire or polymeric filament. In some variations, materials known as Liquid Crystal Polymers (LCP's) may be employed simply because they have extremely high tensile strength and do not easily stretch. This braid allows the  
10 overall diameter of the core wire assembly -- the core wire (260) and the braid (268) -- to be smaller than a core wire having appreciably the same performance. These more complex designs may be useful in situations in which extremely small vasculature must be penetrated to treat the errant thrombus.

As noted elsewhere, the location of the cage assembly should be visible using  
15 fluoroscopy. Various methods have been suggested above. One highly preferred method is shown in Figures 5A and 5B. In essence, at least some of the elements (ribbons or wires) making up the cage assemblies are provided with significant radio-visibility via the placement of a radio-opaque covering on these elements. A metallic coating of a metal having comparatively more visibility, during fluoroscopic use, than  
20 stainless steel is preferred. Such metals are well-known but include gold and members of the Platinum Group of the Periodic Table, e.g., platinum, palladium, rhenium, rhodium, etc. The cage assembly wires or ribbons may be electroplated or otherwise provided with a continuous coating but a thick coating may be had by wrapping the element with a radio-opaque wire or ribbon.

25 Another material suitable for such a service is a structural material filled with a radio-opaque material. For instance, a suitably fine tubing of, e.g., nitinol or

stainless steel, filled with platinum or gold is commercially available. Fort Wayne Metals of Fort Wayne, Indiana sells such a product as a "drawn filled tube."

Figure 5A shows a portion of a cage assembly (300) made up of a woven braid of wires (302) which in turn are tightly wrapped with radio-opaque wires (304). The right side of Figure 5A is a partial cut away showing the wires (302) and the radio-opaque wires (304) in cross-section.

As an example of the device shown in Figure 5A, we have found that nickel/titanium super-elastic alloy wire having a diameter in the range of 0.001" to 0.004" wrapped with 0.0001" to 0.002" diameter platinum wire is suitable and is quite visible.

Figure 5B shows a partial cut-away of a cage assembly (308) constructed of ribbon (310) as described above and further wrapped with radio-opaque wires (312). As was the case with the Figure 5A variation, the right side of the drawing is a partial cut away showing the internal ribbon (310).

The variations of the cage assemblies shown in Figures 5A and 5B may be used in place of any of the cage assemblies found elsewhere herein.

Figures 6A to 6D, show in generic fashion, one method of using the device described herein.

Figure 6A schematically depicts an artery (400) containing a blocking clot (402). Blood flow through artery (400) is severely compromised. A variation of the inventive device (404) similar to that shown in Figures 2A and 2B is shown in the artery (400) in Figure 6B. This variation (404) has a rotatable core wire (406) which is useful as a guidewire. It has a distal woven braided cage assembly (408), a proximal braided cage assembly (410) and a tubular actuator (420). Also shown is the delivery catheter (412) having orifices (414) for perfusion of appropriate medicines.



Many clots such as clot (402) will push the core wire (406) tip aside as the guidewire contacts the clot. Indeed, in most instances, such is preferred. The core wire (406) then slips along the wall of the artery (400) as the device (404) is advanced.

5           Figure 6C shows an appropriate placement of the distal cage assembly (408) past the occlusion (402) and of the proximal cage assembly (410) adjacent to the clot prior to deployment. The proximal cage assembly (410) length assembly is desirably selected so that it overlaps both ends of clot (402).

10           Upon expansion of the cage assemblies (408, 410) to their expanded shape, blood will flow from the arterial to venous end of the clot (402) through the lumen of the open proximal cage assembly (410). Desirably, distal cage assembly (408) has an expanded diameter which approximates the size of the artery (400) lumen. The distal filter assembly (408) is there solely to prevent distal movement of the clot (402) or fragments of the clot. Thrombolytics may be introduced through the orifices (416) to  
15           dissolve clot (402). It is highly desirable that the proximal cage assembly (410) be used to restrain the clot (402) from movement in the vessel (400) so that the thrombolytics may completely dissolve the clot (402) while so restrained. This procedure obviously provides a continuing source of thrombolytics to the clot (402) and further provides a large clot surface area upon which the thrombolytics may  
20           work.

          Alternatively, or in addition to the steps outlined above, clot (402) can be retained or retrieved by the expanded distal cage assembly (408) and removed to another site along with the distal cage assembly (408) and catheter (412).

          It should be apparent that the cage assembly of this invention may be re-  
25           collapsed and re-deployed as necessary to displace or collect emboli as necessary or desired.

The design of such variations is routine and is within the skill of an ordinary designer in this art. Similarly, obvious variants of the invention described above are within the scope of the invention. This and other known or non-critical items have been omitted from the drawings for the sake of brevity and clarity of the invention disclosed here.

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